California Medical Device Recall Information

Recall Name
Hospira Recalls Intravascular Administration Sets Due to a Potential of Puncture

Recall Date	Product Description	Recalling Firm	Recall Reason
Initial: 4/01/13 Updated: 8/13/13	Intravascular Administration Sets	**Hospira, Inc.** Lake Forest, IL	*Potential for the piercing pin on the Hospira Blood Sets to puncture the outer wall of non-ISO-compliant blood bags.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Hospira Blood Sets Listed Products Recalled: • 14200-04-28 Secondary Blood Set; • 14203-04-28 Blood Set; • 14206-04-28 Y-Type Blood Set; • 14207-04-28 Blood Set; • 14210-04-28 Plum Blood Set; • 14211-04-28 Plum Blood Set; • 14212-04-28 Plum Y-Type Blood Set; • 14217-04-28 Y-Type Blood Set • 14219-04-28 Y-Type Blood Set	**CA**, nationwide	Distributed from: July 2011 through February 2013

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm364988.htm